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(21) International Application Number: PCT/US99/05380 (22) International Filing Date: 11 March 1999 (11.03.99) (30) Priority Data: 09/196,848 20 November 1998 (20.11.98) US (71) Applicant: MINNESOTA MINING AND MANUFACTURING COMPANY [US/US]; 3M Center, P.O. Box 33427, Saint Paul, MN 55133-3427 (US). (72) Inventors: KEYES, Michael, P.; P.O. Box 33427, Saint Paul, MN 55133-3427 (US). RADCLIFFE, Marc, D.; P.O. Box 33427, Saint Paul, MN 55133-3427 (US). MARTIN, Steven, J.; P.O. Box 33427, Saint Paul, MN 55133-3427 (US). SNUSTAND, Daniel, C.; P.O. Box 33427, Saint Paul, MN 55133-3427 (US). EPSTEIN, Kenneth, A.; P.O. Box 33427, Saint Paul, MN 55133-3427 (US). (74) Agents: KOKKO, Kent, S. et al.; Minnesota Mining and Manufacturing Company, Office of Intellectual Property Counsel, P.O. Box 33427, Saint Paul, MN 55133-3427 (US).	(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>	

(54) Title: CHIRAL LIQUID CRYSTAL COMPOUNDS HAVING A FLUORINATED TERMINAL PORTION

(57) Abstract

Novel liquid crystal compounds are provided which exhibit less temperature dependent switching properties, for the reliable and consistent operation of liquid crystal devices. The liquid crystal compounds comprise (a) an achiral fluorochemical terminal portion that comprises a terminal fluoroalkyl, fluoroether, perfluoroalkyl, or perfluoroether group; a chiral terminal portion comprising a saturated or unsaturated, chiral hydrocarbon or chiral hydrocarbon ether group comprising a chiral center; and a central core connecting the terminal portions; said chiral terminal portion having at least three in-chain atoms (an "extended group") between said chiral center of the chiral terminal portion and said central core.

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More recently, antiferroelectric liquid crystals (AFLC), another class of tilted chiral smectic liquid crystals, have been developed. Antiferroelectric liquid crystals are

switchable in a chiral smectic C_A phase (SC_A* phase) in addition to the tilted chiral smectic C phase (SC* phase) used in ferroelectric liquid crystal devices.

One of the most important characteristics of a liquid crystal display device is its response time, i.e., the time required for the device to switch from the on (light) state to the off (dark) state. Especially important for practical operation of larger devices is the response time with respect to temperature. In such devices, temperature non-uniformities may adversely effect the performance and require some form of compensation, unless the switching speed is largely independent of temperature. In a ferroelectric or anti-ferroelectric device, response time τ_{electric} is proportional to the rotational viscosity (η) of the liquid crystal compound(s) contained within the device and is inversely proportional to their polarization (P_s) and to the applied electric field (E) according to the following formula:

$\tau_{\text{electric}} = \eta / P_s E$. Thus, response time can be reduced by using compound(s) having high polarizations or low viscosities, and such compounds are greatly desired in the art. In addition, compounds with polarizations that rapidly increase with decreasing temperature can lead to temperature independence or reduced temperature dependence of switching.

In the passive addressing of liquid crystal compounds exhibiting a spontaneous polarization, however, low polarization mixtures may be preferred for the practical operation of a liquid crystal device. Polarization reversal fields are larger for higher polarization mixtures, and polarization reversal fields cause switching or partial switching back to a material's original director alignment. This results in loss of the bistability that is crucial to the passive-matrix driving of ferroelectric liquid crystal devices.

Another potential disadvantage of using high polarization mixtures is the partial switching of their director alignment in response to non-switching (secondary) signals in a driving waveform. This continued response or fluctuation of the director causes a large decrease in the contrast ratio of a ferroelectric liquid crystal device.

There remains a need in the art for liquid crystal materials having fast response times, ideally possessing broad smectic temperature ranges to enable operation of the device over a broad range of temperatures, or should be capable of combination with other liquid crystal compounds having different smectic temperature ranges without adversely

affecting the smectic phase behavior of the base mixture. There further remains a need in the art for materials which have low polarization values, and which can provide reduced temperature dependence of the response time. Further, there remains a need in the art for novel liquid crystal materials which exhibit tristable switching.

5

Summary of the Invention

Briefly, in one aspect, this invention provides fluorine-containing, chiral liquid crystal compounds having smectic mesophases or latent smectic mesophases.

(Compounds having latent smectic mesophases are those which by themselves do not exhibit a smectic mesophase, but which, when in admixture with compounds having smectic mesophases or with other compounds having latent smectic mesophases, develop smectic mesophases under appropriate conditions.)

The chiral liquid crystal compounds of the invention comprise (a) an achiral fluorochemical terminal portion which comprises a terminal fluoroalkyl, fluoroether, perfluoroalkyl, or perfluoroether group; (b) a chiral terminal portion comprising a saturated or unsaturated, chiral hydrocarbon or chiral hydrocarbon ether group comprising a chiral center; and (c) a central core connecting the terminal portions; the chiral terminal portion having at least three in-chain atoms (an "extended group") between said chiral center of the chiral terminal portion and the central core.

Surprisingly, in comparison with similar compounds having fewer than three in-chain atoms between a chiral center of the chiral terminal portion and the central core, the compounds of the invention provide comparable electrooptic response speeds in spite of their lower measured polarization values. These lower polarization values, in combination with broad mesogenic temperature ranges, enable the utilization of liquid crystal mixtures that contain up to 100% of the chiral (optically active) compounds of the invention. In general, mixtures containing a high concentration of the compounds of this invention exhibit less temperature dependent switching properties, which is important for the reliable and consistent operation of liquid crystal devices.

Furthermore, the use of high concentrations of liquid crystal compounds having low polarizations also provides a decrease (relative to the use of low concentrations of compounds having high polarizations) in the partial switching response of the resulting

compositions to non-switching (secondary) signals in the driving waveform that is commonly used in the passive addressing of liquid crystal devices. Such a decrease in this response is critical for improving the contrast of a device.

5 In another aspect, this invention provides liquid crystal display devices utilizing both the liquid crystal compounds of the invention and mixtures containing the liquid crystal compounds of the invention. In yet another aspect, the invention provides liquid crystal compounds that exhibit tristable switching, and liquid crystal display devices of the invention that can provide tristable switching, gradation display (grey scale), threshold control, hysteresis control, and fast response times. The devices can be either passive or
10 active matrix devices.

In yet another aspect, this invention provides a process for reducing the temperature dependence of the switching speed of a liquid crystal mixture by combining at least one liquid crystal compound of this invention with at least one other compound liquid crystal composition comprising at least one smectic or latent smectic liquid crystal
15 compound having a temperature dependent switching with slope typically more negative than -0.02 , preferably more negative than -0.03 .

These and other features, aspects, and advantages of the present invention will become better understood with regard to the following description, examples and accompanying Figures.

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Brief Description of the Figures

Figures 1 and 2 show the temperature dependence of the polarization (P_s) of the compounds of the invention.

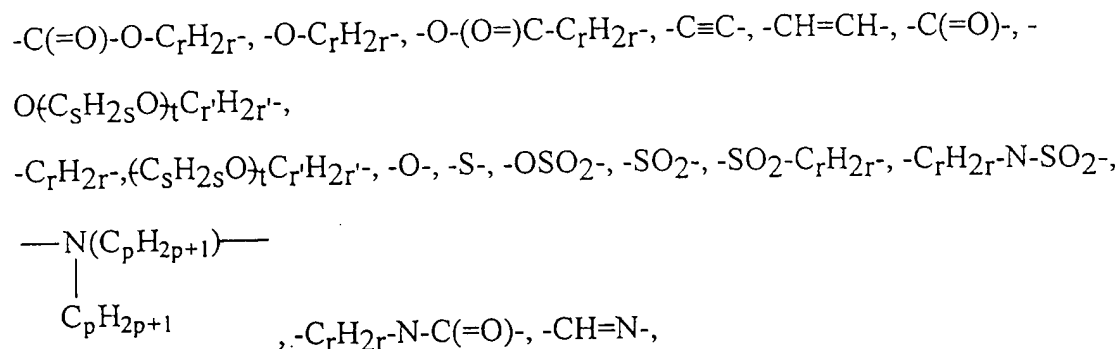
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Figure 3 show the temperature dependence of the switching speed τ_{electric} of the compounds of the invention.

Figures 4 and 5 show tristable switching behavior of selected compounds of the
30 invention.

Detailed Description of the Invention

The chiral terminal portion of the compounds of the invention can be represented by the formula $-D-R^*$, where R^* is a saturated or unsaturated, chiral hydrocarbon or hydrocarbon ether moiety containing at least one chiral center (asymmetric carbon atom); and D is non-directionally selected from the group consisting of a covalent bond,



10 and combinations thereof, and where r and r' are independently integers of 0 to about 20, s is independently an integer of 1 to about 10 for each $(C_5H_2SO)_t$, t is an integer of 1 to about 6, and p is an integer of 0 to about 4; with the proviso that at least one chiral center of R^* is spaced from the central core by at least 3 in-chain atoms. Preferably D of $-D-R^*$ is $-O-C_rH_{2r}-$ or $-C_rH_{2r}-$.

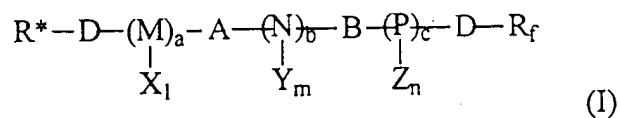
15 The achiral fluorochemical terminal portion of the compounds of the invention can be represented by the formula $-D-R_f$, where D is as previously defined and where R_f is an achiral fluoroalkyl, perfluoroalkyl, fluoroether, or perfluoroether group. Preferably, R_f is perfluoroalkyl or perfluoroether; more preferably, R_f is perfluoroether, as the perfluoroether-containing compounds of the invention exhibit, e.g., a broad smectic C mesophase, good compatibility with other smectic C compounds, and advantageous layer spacing behavior. When the R_f group of the fluorochemical terminal portion is perfluoroalkyl or perfluoroether, it can contain small amounts of residual carbon-bonded hydrogen atoms but is preferably completely fluorinated. Preferably D of $-D-R_f$ is $-O-C_rH_{2r}-$, $-O(C_5H_2SO)_tC_{r'}H_{2r'}-$, $-C_rH_{2r}-$, or $(C_5H_2SO)_tC_{r'}H_{2r'}-$.

25 In general, the compounds of this invention have a central core comprised of at least one or two rings independently selected from the group consisting of aromatic, heteroaromatic, alicyclic, substituted aromatic, substituted heteroaromatic, and substituted alicyclic rings, the rings being connected one with another by a covalent bond or by

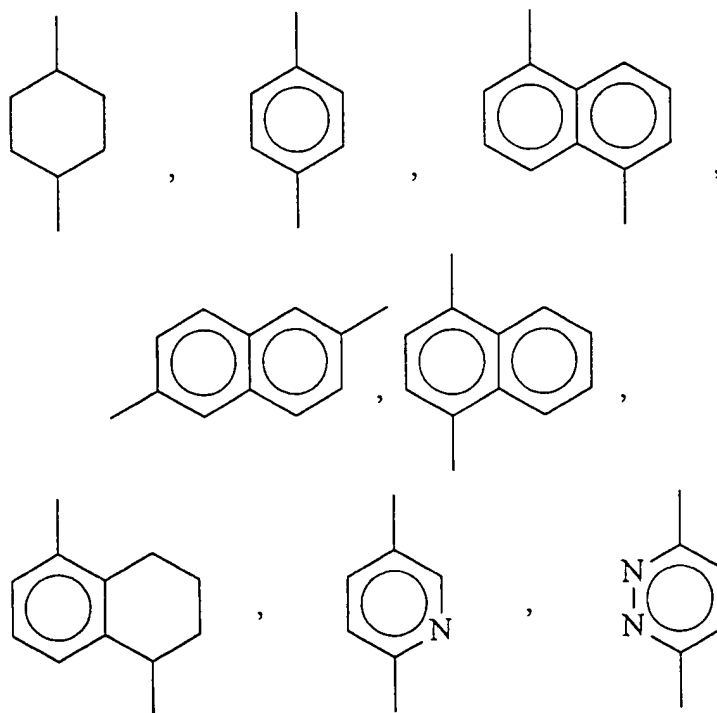
chemical groups selected from the group consisting of -COO-, -COS-, -HC=N-, -CH=CH-,
 -C≡C-, and -COSe-. The rings can be fused or non-fused. The heteroatoms within the
 heteroaromatic rings comprise at least one atom selected from the group consisting of
 nitrogen, oxygen, and sulfur. Non-adjacent ring carbon atoms in the alicyclic rings can be
 5 substituted by nitrogen, oxygen, or sulfur atoms. When the ring(s) are aromatic,
 heteroaromatic, substituted aromatic, or substituted heteroaromatic, the non-fused rings of
 the core are preferably no more than about two in number.

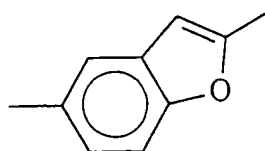
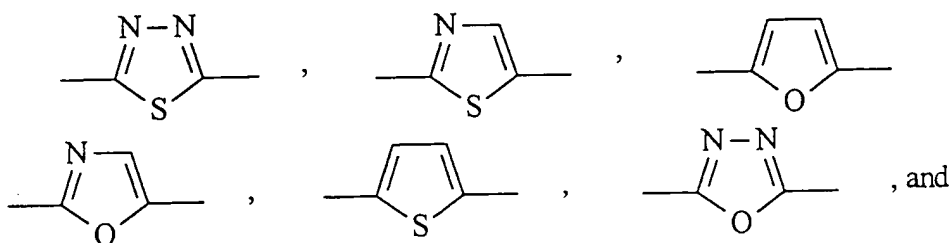
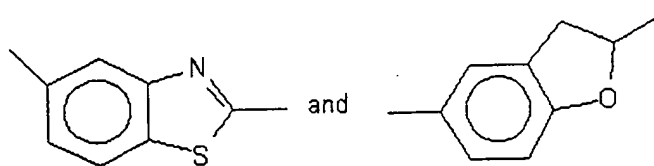
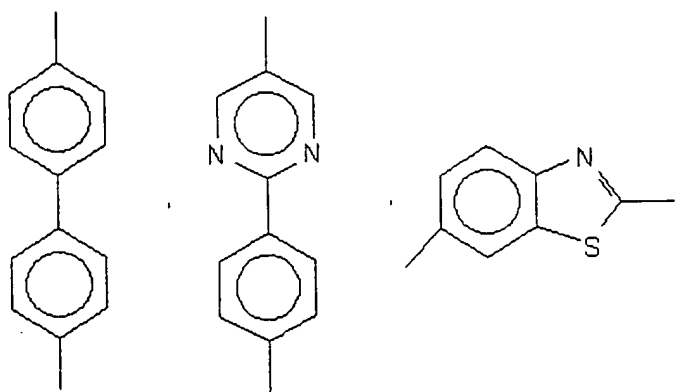
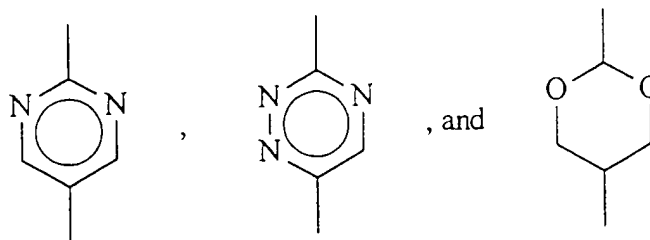
The chiral liquid crystal compounds of the invention exhibit exceptionally wide
 mesomorphic temperature ranges. When used in electrooptical devices, the compounds
 10 provide fast response times upon application of an electric field over broad temperature
 ranges. This makes them extremely useful in the preparation of mixtures that can operate
 in their active mesomorphic phase from around -30°C to around 70°C.

A class of the above-described liquid crystal compounds of the present invention
 can be represented by the general formula I:



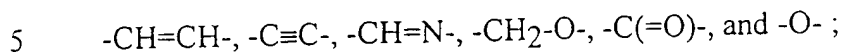
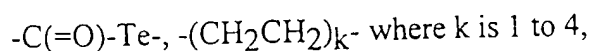
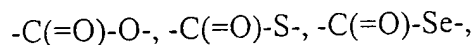
15 where M, N, and P are each independently selected from the group consisting of





a, b, and c are each independently zero or an integer of from 1 to 3, with the proviso that the sum of $a + b + c$ be at least 1 (and preferably no greater than 2);

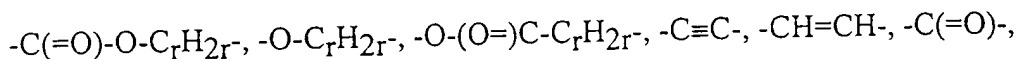
each A and B are non-directionally and independently selected from the group consisting of a covalent bond,



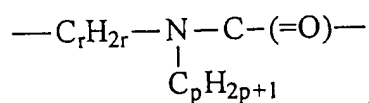
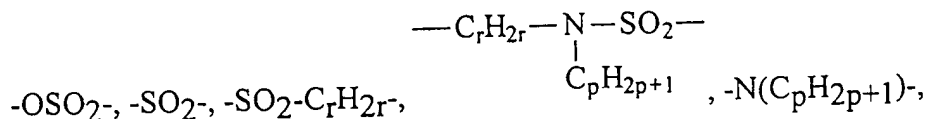
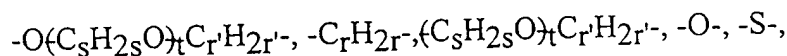
each X, Y, and Z are independently selected from the group consisting of -H, -Cl, -F, -Br, -I, -OH, -OCH₃, -CH₃, -CF₃, -OCF₃, -CN, and -NO₂;

10 each l, m, and n are independently zero or an integer of 1 to 4;

each D is non-directionally and independently selected from the group consisting of a covalent bond,



15



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where one or more hydrogen atoms can optionally be replaced with fluorine, and where r and r' are independently integers of 0 to about 20, s is independently an integer of 1 to about 10 for each (C₅H₂S₂O), t is an integer of 1 to about 6, and p is an integer of 0 to about 4;

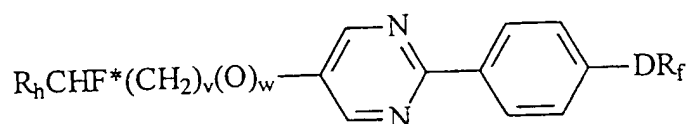
25 R* is a chiral hydrocarbon or hydrocarbon ether moiety containing at least one chiral center (asymmetric carbon atom) with the proviso that at least one chiral center of R* is spaced from the central core by at least 3 in-chain atoms;
R_f is fluoroalkyl, perfluoroalkyl, fluoroether, or perfluoroether, preferably perfluoroalkyl or perfluoroether. More preferably, R_f is perfluoroether.

In defining R_f , particularly preferred perfluoroalkyl groups are those which can be represented by the formula $-C_qF_{2q}X'$, where q is an integer of 1 to about 20 (and, preferably, is at least about 5) and X' is hydrogen or fluorine. Particularly preferred perfluoroether groups are those which can be represented by the formula -

5 $(C_xF_{2x}O)_zC_yF_{2y+1}$, where x is independently an integer of 1 to about 12 for each $(C_xF_{2x}O)$, y is an integer of 1 to about 12, and z is an integer of 1 to about 10. Preferably, the perfluoroether group is linear, x is independently an integer of 1 to about 6 for each $(C_xF_{2x}O)$, y is an integer of 1 to about 6, and z is an integer of 1 to about 6.

A preferred subclass of the above-described chiral compounds of the invention can be represented by the formula:

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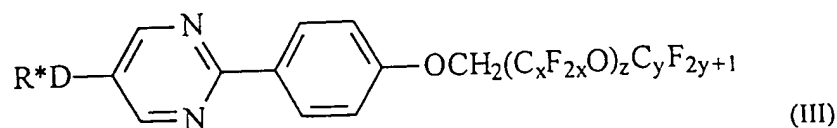


where D and R_f are as previously defined, R_h is a linear or branched alkyl group of 1 to about 10 carbon atoms, $-CHF^*$ represents a chiral center of either (S) or (R) configuration, v is an integer from 2 to about 10, w is an integer of 0 or 1, with the proviso that $v+w$ is at least 3, where one or more non-adjacent $-CH_2-$ groups can be replaced by an oxygen atom and where one or more ring hydrogen atoms can be replaced by fluorine atoms.

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Another preferred subclass of the above-described extended chiral compounds of the invention may be represented by the formula:

20



where R^* , D , x , y and z are as previously defined, and where one or more ring hydrogen atoms may be replaced by fluorine atoms.

25 Surprisingly, many of the chiral liquid crystal compounds of Formula III have been found to exhibit tristable switching properties, making them useful as liquid crystal

materials in both passive and active-matrix tristable devices having tristable switching, gradation display (grey scale), threshold control, hysteresis control, and fast response times.

Devices employing antiferroelectric liquid crystals (AFLC) have been described by Chandani et. al (Japan J. of Applied Physics 27(5), L729-732 (1988).). Antiferroelectric liquid crystals used in these devices exhibit three stable states: two stable states under the influence of an electric field and a third antiferroelectric state in the absence of an electric field. Antiferroelectric liquid crystals are characterized by having a distinct threshold and a double hysteresis that allows for a memory effect in either of the driven states.

Antiferroelectric liquid crystals can be easily switched and provide devices that have few defects and that allow for the recovery of alignment.

In an AFLC device, with no applied electric field, an AFLC composition has a layered structure comprising many smectic layers, with the molecules of each layer being tilted in a direction opposite to those of the adjacent layer such that the liquid crystal composition has no net polarization. The alternating molecular director also results in a uniform optical axis parallel to the layer normal of the smectic layers. When placed between a pair of crossed polarizers such that one of the polarization axes of the polarizers is aligned with the uniform optical axis of the composition, the device exhibits a dark state. When an electric field is applied, the liquid crystal orients to align the spontaneous polarization with the electric field, resulting in one of two bright states, depending on the polarity of the electric field. Tristable switching behavior has also been observed for twisted ferroelectric and deformed helix devices.

The tristable liquid crystal display device of the invention comprises (a) first and second opposed substrates, at least one bearing an alignment coating and each bearing at least one electrode so as to define one or a plurality of pixels; (b) a tilted smectic or induced tilted smectic liquid crystal composition comprising at least one compound of Formula III disposed between the substrates; and (c) a pair of orthogonally disposed polarizers, each having a polarization (or light transmission) axis, one polarization axis being aligned with the zero field optical axis of a tilted smectic or induced tilted smectic mesophase of the liquid crystal composition; wherein the substrates are disposed so as to provide an alignment of the liquid crystal composition.

The substrates can comprise any of the materials known in the art to be useful as substrates for liquid crystal display devices, e.g., glass or plastic. The electrodes can be of any electrically conductive material, e.g., indium tin oxide (ITO), and can be applied to the surface of the substrates by any of the methods known in the art. Thus, the substrates can be coated with a film of SnO_2 , InO_3 , or ITO to form electrodes.

The substrates and the electrodes bear alignment coatings, which can comprise any of the useful alignment compositions known in the art. The two coatings can be the same or different. Materials suitable for use in preparing alignment coatings include polyvinyl alcohol, polyimide, polyamide-imide, polyester, polyamide, polyester-imide, polyparaxylylene, polycarbonate, polyvinyl acetal, polyvinyl chloride, polystyrene, polysiloxane, cellulose resin, melamine resin, urea resin, acrylic resin, and the like, and mixtures thereof. The surface of the alignment coatings can be subjected to a prescribed (uniaxial) alignment treatment as desired, for example, by rubbing the surface with a fibrous material such as velvet, cloth, or paper.

Substrates can be subjected to different alignment treatments in order to appropriately control an alignment state, particularly an initial alignment state. For example, one of the substrates can be provided with a rubbing-treated alignment coating, and the other can be provided with an alignment coating that is not subjected to rubbing and/or that comprises a composition that is different from that of the rubbing-treated alignment coating.

The liquid crystal display device of the invention preferably has at least one alignment coating of sufficient thickness to optimize the tristable switching properties of the device. The coating preferably has a thickness of from about 50 to about 5000 angstroms, more preferably from about 50 to about 2500 angstroms, in order to provide good driving characteristics, high reliability, and driving stability in a wide temperature range.

Preferably, one of the alignment coatings of the device of the invention comprises a polyimide or polyamide (e.g., nylon) film. The film can generally be prepared by applying a polyamic acid (polyimide precursor) or polyamide solution to the surface of the substrate, heating the applied coating layer, and then subjecting the resulting alignment coating to a rubbing treatment. If desired, the device of the invention can utilize alignment

treatments such as those described, e.g., in EP 755993 (Canon) and U.S. Patent No. 5,377,033 (Radcliffe). The device of the invention can optionally further comprise an insulating layer.

Substrates bearing said alignment coatings are separated by spacers at a fixed distance (the "cell gap") that, along with the alignment coatings, allows for the alignment of a liquid crystal composition that is contained in the resulting space between the substrates. The cell gap can generally be up to about 10 μ , preferably from about 0.5 to about 5 μ , in order for the device to exhibit tristable switching.

On the outer surfaces of substrates are affixed orthogonally disposed polarizers, each having a polarization (or light transmission) axis. The polarizers can be of any design and material known in the art to be useful in liquid crystal display devices. The polarization axis of one of the polarizers is aligned with the zero field optical axis of a tilted or induced tilted smectic mesophase of the liquid crystal composition.

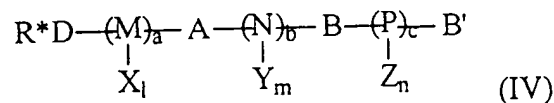
The device of the invention containing the above-described liquid crystal composition can be driven by either a passive or an active matrix display. A typical liquid crystal display consists, e.g., of two polarizers, two transparent substrates, switching elements or electrodes to define pixels, and driver Integrated Circuits (ICs) to address rows and columns of pixels. The rows and columns can be constructed out of strips of conductive material.

In a passive matrix display, pixel-matrix elements can be defined by the intersection areas of rows and columns of a transparent conductor material, e.g., indium tin oxide (ITO), on the inner surfaces of two opposed substrates. To switch or address a liquid crystal composition disposed between these pixel-matrix elements, charge is applied to the appropriate row and column in order to change the orientation of the liquid crystal material (i.e., to change from a dark pixel to a bright pixel). Passive matrix displays have been described, for example, by Peter J. Collings in Liquid Crystals: Nature's Delicate Phase of Matter, pp. 100-103, Princeton University Press, Princeton, N.J (1990) and by Peter J. Collings and Michael Hird in Introduction to Liquid Crystals Chemistry and Physics, pp. 271-285, Taylor and Francis Ltd., London (1997).

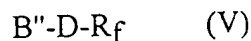
In order to improve on the limitations of passive matrix displays (such as crosstalk caused by driving waveforms), active matrix displays have been developed in the art.

Active matrix displays typically have thin film transistors (TFTs) or diode arrays on a glass substrate, which address each pixel element. The TFT can comprise amorphous silicon (α -Si) or polycrystalline silicon (p-Si) or can comprise a single crystal semiconductor device such as a CMOS (Complementary Metal Oxide Semiconductor) silicon-based device. The TFT electrically isolates one pixel element from the others in the display and eliminates the problem of partially active pixels. TFT can simply be considered a switch; when selected (on), it allows charge to flow through it, and, when off, it acts as a barrier that prevents or at least restricts the flow of charge. When a row of TFTs is addressed, gate lines are active, and the "switch" is turned on, allowing charge to flow from the columns into the pixels and to set the image for the frame cycle. Once a row has been addressed, the gate line is reverse biased (the switch is turned off) to insure that no charge can pass from the columns into the pixel elements. Thus, the pixel is now isolated as the rest of the display is addressed. Active matrix arrays have been described, for example, by Collings *supra* and in U.S. Patent No. 5,631,752 (Tonaka).

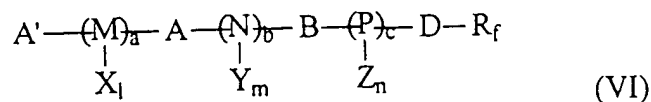
The fluorine-containing liquid crystal compounds of the invention can be prepared by a process comprising the steps of (a) mixing at least one compound represented by the formula



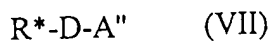
with at least one compound represented by the formula



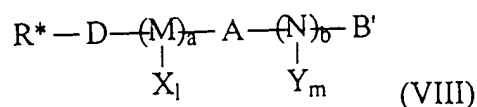
or mixing at least one compound represented by the formula



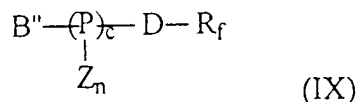
with at least one compound represented by the formula



or mixing at least one compound represented by the formula



with at least one compound represented by the formula



where M, N, P, a, b, c, A, B, X, Y, Z, l, m, n, D, R*, and R_f are as defined above for formula I; x is an integer of 0 or 1; and each A', A'', B', and B'' are independently selected from the group consisting of -H, -Cl, -Br, -I, -OH, -COOH, -CH(CH₂OH)₂, -SH, -SeH, -TeH, -NH₂, -COCl, -CHO, -OSO₂R_f'', -OSO₂CH₃, -C≡CH, dialkyl borane, -CH=CH₂, -NH(C=O)OC_qH_{2q+1}, -NCO, -OSO₂-cyclo(C₆H₄)-CH₃, -CH₂COOH, and -CH(C(O)O-C_qH_{2q+1})₂, where R_f''' is a perfluoroalkyl group having from 1 to about 10 carbon atoms and q is an integer of 0 to about 20, and with the proviso that R*-D-A'' can enter into an addition or condensation reaction with A' and that B''-D-R_f can enter into an addition or condensation reaction with B';

and (b) allowing compounds IV and V, compounds VI and VII, or compounds VIII and IX to react, optionally in the presence of suitable coupling agent(s), i.e., reagent(s) which effect coupling. For Formula VII A'' is preferably -OH (more preferably -CH=CH₂), and -D-R_f is preferably -OCH₂-R_f.

The compounds of the invention are useful in admixture with themselves or with other chiral or achiral liquid crystal compounds (as dopants or as the major components), for electrooptical display applications. The compounds have a number of desirable properties when used in admixture with themselves or with other liquid crystal compounds, preferably compounds having fluorinated terminal portions such as those compounds disclosed, for example, in U.S. Pat. Nos. 4,886,619 (Janulis), 5,082,587 (Janulis), 5,254,707 (Janulis), 5,262,082 (Janulis et al.), 5,377,033 (Radcliffe), 5,399,291 (Janulis et al.), 5,399,701 (Janulis), 5,437,812 (Janulis et al.), 5,474,705 (Janulis et al.), 5,482,650 (Janulis et al.), 5,658,491 (Kistner et al.), 5,702,637 (Johnson et al.), and U.S.S.N 08/998,400 filed 12/24/97 (Attorney's docket number 53589USA5A).

For example, the compounds of the invention when admixed with such preferred liquid crystal compounds show excellent compatibility, show a beneficial effect or only a minimal negative effect on the smectic C temperature range of the resulting mixtures (even when present at high concentrations), and provide tilted smectic mixtures (ferroelectric, antiferroelectric, or twist) having fast electrical response times. Mixtures containing the compounds exhibit favorable alignment, switching, response to an electric field, temperature dependence of response speed, temperature dependence of polarization, contrast, layer structure, and mesomorphic temperature ranges. Compounds of the invention can also be used to optimize mixture properties such as tilt angle, memory angle, spontaneous polarization and its temperature dependence, liquid crystal temperature range, switching behavior, birefringence, and the temperature dependence of layer spacing.

Most of the compounds of the present invention have enhanced smectic mesophases. Mixtures of the compounds of the invention with other liquid crystal materials can be formulated to provide desired transition temperatures and broad mesophase temperature ranges. Such mixtures preferably contain compounds having fluorinated terminal portions, such as those compounds described, for example, in U.S. Pat. Nos. 4,886,619 (Janulis), 5,082,587 (Janulis), and, most preferably, 5,262,082 (Janulis et al.). The liquid crystal compounds of the invention can also be used to prepare ferroelectric liquid crystal devices such as, e.g., those described in U.S. Patent Nos. 5,377,033 (Radcliffe), 5,417,883 (Epstein et al.) and 5,641,427 (Shinjo) and in EP 769542 and EP 769543.

The compounds of this invention in admixture with other chiral or achiral liquid crystal compounds may exhibit chiral smectic liquid crystal behavior. Furthermore, many of the perfluoroether group-containing liquid crystal compounds of the invention when used alone or when mixed with other liquid crystal compounds of the invention or with achiral, fluorine-containing liquid crystal compounds (preferably, the perfluoroether group-containing liquid crystal compounds described in U.S. Pat. No. 5,262,082 (Janulis et al.)) exhibit a reduced temperature dependence of the smectic interlayer spacing. This property enables the spontaneous generation of a bookshelf type layer structure, which is ideal for a tilted smectic liquid crystal device. In general, the compounds of the invention

exhibit maintenance or expansion of the smectic C layer spacing with decreasing temperature.

In the process of this invention, liquid crystal mixtures can be obtained having reduced temperature dependence of switching. With many liquid crystal mixtures, the switching speed, τ_{electric} increases with decreasing temperature, resulting in unacceptably slow switching speeds at operating temperatures of a device containing the liquid crystal mixture. Frequently, liquid crystal compounds having either high polarization (P_S) or low rotational viscosity (η) are added to a mixture in order to reduce the response time, but such methods do not reduce the temperature dependence of the response time.

The present invention provides a process for reducing the temperature dependence of the response time of a liquid crystal mixture, which comprises the step of combining:

(a) at least one compound of the invention which comprises (i) an achiral fluorochemical terminal portion which comprises a terminal fluoroalkyl, fluoroether, perfluoroalkyl, or perfluoroether group; (ii) a chiral terminal portion comprising a saturated or unsaturated chiral hydrocarbon or chiral hydrocarbon ether group comprising a chiral center; and (iii) a central core connecting the terminal portions; the chiral terminal portion having at least three in-chain atoms between the chiral center and the central core; with

(b) a liquid crystal composition comprising at least one smectic or latent smectic liquid crystal compound exhibiting temperature dependent switching (when placed in a device)).

The compound(s) of composition (b), typically exhibit a plot of $\ln \tau_{\text{electric}}$ vs. temperature, having a slope (the change of \ln (natural log) of τ_{electric} with ΔT) typically more negative than -0.02, preferably more negative than -0.03. Especially suitable compounds for use in component (b) are fluorine containing, smectic or latent smectic liquid crystal compounds (preferably compounds having fluorinated terminal portions such as those compounds described, for example, in U.S. Pat. Nos. 4,886,619 (Janulis), 5,082,587 (Janulis), 5,254,707 (Janulis), 5,262,082 (Janulis et al.), 5,377,033 (Radcliffe), 5,399,291 (Janulis et al.), 5,399,701 (Janulis), 5,437,812 (Janulis et al.), 5,474,705 (Janulis et al.), 5,482,650 (Janulis et al.) 5,658,491 (Kistner et

WO 00/51210

al.), 5,702,637 (Johnson et al.), and U.S.S.N 08/998,400 filed 12/24/97 (Attorney's docket number 53589USA5A).

The process of the invention can be carried out by combining components (a) and (b). The combining or mixing of the components can be effected by introducing the components to a vessel, generally with simultaneous and or subsequent agitation or stirring, e.g., roller mixing. The vessel can be either an open or a closed vessel of a size which is sufficient to hold both compositions while allowing room for mixing. The components can be formed prior to combination with each other, or alternatively, one of more of the compounds of either can be combined with one or more of the compounds of the other prior to addition of the remaining compounds. Any order and manner of combination of the compounds of the two components is acceptable. The resulting combination is preferably agitated or stirred sufficiently so that a homogeneous mixture is achieved. This is preferably facilitated by applying sufficient heat to melt the combination, or by dissolving the combination in a solvent, e.g., a polar aprotic solvent, with subsequent solvent removal, e.g., by evaporation.

These components (a) and (b) are mixed in ratios so as to balance or counteract the large negative slope of component (b) with the less negative or positive-tending slope of component (a). The liquid crystal compounds to be utilized in the process can be selected based upon the sign and magnitude of the $\Delta(\ln \tau_{\text{electric}}) / \Delta T$ slope (or, in the case of latent smectic liquid crystal compounds, the sign and magnitude of the slope of mixtures containing the latent compounds(s), which can be determined by measuring the switching speed τ_{electric} at the desired temperatures as described below in the Examples. In general, composition (b) (generally having a more negative τ_{electric} slope than -0.02) can be combined with composition (a) (generally having a less negative τ_{electric} slope than -0.03 to -0.02 or even positive τ_{electric} slope) to obtain a combination having a desired intermediate $(\ln \tau_{\text{electric}}) / \Delta T$ slope associated with a reduced temperature dependency of the switching speed.

Preferably, component (a) is utilized in amounts such that the resulting combination has a tau slope between -0.03 and 0.04 (more preferably, between -0.02 and 0.02 ; most preferably, between -0.01 and 0.01). However, in some cases a tau slope

outside of these ranges may be desirable for a particular purpose and can be achieved through an iterative process of combining components (a) and (b) in varying ratios and measuring the net tau slope of the resulting combinations.

Another advantage of using the materials of this invention in the formulation of liquid crystal mixtures is the low birefringence which can be obtained. The low birefringence of the liquid crystal compounds of the invention (relative to their non-fluorine-containing analogues) allows the fabrication of devices with larger device spacings. Light transmission through, e.g., a surface-stabilized ferroelectric device (as described in U.S. Patent Nos. 4,367,924 (Clark et al.) and 5,377,033 (Radcliffe) with two polarizers is represented by the following equation:

$$I = I_0 (\sin^2(4\Theta)) (\sin^2(\pi\Delta nd/\lambda))$$

where I_0 = transmission through parallel polarizers

- Θ = material tilt angle
- Δn = liquid crystal birefringence
- d = device spacing
- λ = wavelength of light used

To maximize the transmission in a bistable device, both $\sin^2(4\Theta)$ and $\sin^2(\pi\Delta nd/\lambda)$ must be at maximum. This occurs when each term equals one. The first term is a maximum when the tilt angle equals 22.5° . This is a function of the liquid crystal and is constant for a given material at a given temperature. The second term is maximum when $\Delta nd = \lambda/2$. This demonstrates the importance of the low birefringence of the materials of this invention. Low birefringence allows a larger device thickness, d , for a given wavelength of light. Thus, a larger device spacing is possible while still maximizing transmission, allowing easier device construction.

For a tristable device the following equation is used:

$$I = I_0 (\sin^2(2\Theta)) (\sin^2(\pi\Delta nd/\lambda))$$

To maximize the transmission in a tristable device, both $\sin^2(2\Theta)$ and $\sin^2(\pi\Delta nd/\lambda)$ must be at maximum. This occurs when each term equals one. The first term is a maximum when the tilt angle equals 45° .

Objects and advantages of this invention are further illustrated by the following examples, but the particular materials and amounts thereof recited in these examples, as well as other conditions and details, should not be construed to unduly limit this invention.

EXAMPLES

In the following examples, all temperatures are in degrees Celsius and all parts and percentages are by weight unless indicated otherwise. Commercially available materials were chemically transformed by reaction pathways well-known to those skilled in the art and detailed in the examples. Chemical transformations were comprised of acylation, esterification, etherification, alkylation, and combinations thereof using fluorine-containing and non-fluorine-containing reactants to provide the precursor compounds, which, in turn, were caused to react together to yield the chiral, fluorine-containing liquid crystal compounds of this invention.

Compounds prepared in the various examples of this invention were characterized by their melting or boiling point, and structures were confirmed by using at least one of the following methods of analysis: chromatography; ^{13}C -, ^1H -, and ^{19}F -NMR; and infrared and mass spectroscopies.

The 5-alkyl-2-(4-hydroxyphenyl) pyrimidines used in the examples were prepared essentially as described by Zschke and Stolle in "Synthese niedrigschmelzender Kristallin-Flüssiger Heterocyclen; 5-n-Alkyl-2-[4-n-alkanoyloxy-phenyl]pyrimidine," Z.Chem. 15, 441-3 (1975). (S)- and (R)-2-fluoro-decyl-p-toluenesulfonate were prepared essentially as described by Nohira et al. in Mol. Cryst. Liq. Cryst. 180B, 379 (1990). Fluorinated alcohols were prepared essentially as described in U.S. Patent No. 5,262,082 (Janulis et al.) by sodium borohydride reduction of the corresponding perfluorinated acids (or derivatives), which had been prepared by electrochemical fluorination (ECF) or by direct fluorination (using elemental fluorine) of the corresponding hydrocarbon acids (or derivatives). See, e.g., the description of ECF given in U.S. Patent No. 2,519,983

(Simons). Direct fluorination is described, e.g., in U.S. Patent No. 5,362,919 (Costello et al.).

5 Example 1

Preparation of 5-Fluoro-1-methylsulfonyl heptane

(R)-6-benzyloxyhex-1-ene oxide was prepared by copper catalyzed addition of 3-
10 benzyloxypropyl-1-magnesium bromide to R-epichlorohydrin to form R-1-chloro-6-benzyloxyhexan-2-ol which was then dehydrated under basic conditions to R-6-benzyloxyhex-1-ene oxide. This epoxide was then treated with methyl lithium (1.5 M in diethyl ether) in the presence of dilithiotetrachlorocuprate (0.02 molar equivalents, 0.1M in tetrahydrofuran) to yield S-1-benzyloxy-5-hydroxyheptane.

15 A dry one liter flask was equipped with an overhead stirrer, an addition funnel, and a thermocouple. S-1-benzyloxy-5-hydroxyheptane (11 g, 49.5 mmol) and 140 ml toluene were added to the flask under positive nitrogen pressure and cooled to -15°C. Perfluorobutanesulfonyl fluoride (26.3g, 87.1 mmol) was then added to the cooled mixture. After about 5 minutes 1,8-diazobicyclo[5.4.0]undec-7-ene (DBU, 14 g, 92.1
20 mmol) was added at such a rate that the temperature did not rise above 5 °C. After complete addition, the cooling bath was removed and the reaction mixture was stirred for one hour followed by addition of toluene (140 ml), then water (100 ml). The resultant upper toluene phase was washed with 100 ml 7% HCl, then with 2X 100 ml water. The toluene phase was evaporated under reduced pressure to yield 11.3 g of gold-colored
25 liquid which was shown to be 84% of the desired R-1-benzyloxy-5-fluoroheptane and 13% olefin by gas chromatography. The crude product was further purified by chromatography on silica gel eluting with 40:1 v/v hexane/ ethyl acetate to give 6.05 g of clear liquid; ¹H-nmr showed this material to be 95% desired compound, 0.4% RCH₂F, 1.4% RCH₂CF₂CF₂CF₂F and 3% ethyl ether (mole %).

30 The benzyl protecting group was removed by hydrogenation using 10% Pd on carbon. The resulting alcohol was treated with methanesulfonyl chloride in the presence

of triethylamine to generate the desired intermediate R-5-Fluoro-1-methylsulfonyl pentane.

5 **Preparation of 5-hydroxy-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine**

To a 1 liter round bottom flask fitted with an addition funnel, thermocouple and a mechanical stirrer was added 5-benzyloxy-2-[4-hydroxyphenyl]pyrimidine (20g, .072 mol), perfluorobutanesulfonyl fluoride (26.0, 0.086 mol) and t-butyl methyl ether (204 ml).
10 The solution was cooled to 15°C, and DBU (13.1g, 0.086mol) was added through an addition funnel at a rate sufficient to maintain the reaction temperature below 30°C. The reaction was stirred for 2 hours at room temperature and then washed with water, 10% HCl and brine. The resulting organic layer was separated and concentrated. The product was recrystallized from ethanol to give 5-benzyloxy-2-[4-
15 perfluorobutanesulfonylphenyl]pyrimidine.

To a 250 ml round bottom flask fitted with a magnetic stir bar and thermocouple was added 9-borabicyclo[3.3.1]nonane (9-BBN, 53 ml of 0.5M soln in THF). The solution was stirred under a nitrogen atmosphere then cooled to 5°C. 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene (7.8g, 0.022mol, prepared from 1-bromohex-6-ene and 3-
20 pentafluoroethoxy-2,2,3,3-tetrafluoropropanol) was added, and the mixture was stirred at room temperature for 4 hours. 5-benzyloxy-2-[4-perfluorobutanesulfonylphenyl]pyrimidine (10g, 0.0195mol), dichlorobis(triphenylphosphine) palladium II (0.27g, 0.0004mol), and sodium hydroxide (1.56g, 0.039mol) were added to the reaction mixture. The reaction was heated to 50°C
25 for 2 hours, cooled to room temperature, diluted with toluene (100ml) and washed with water. The resulting organic phase was concentrated under reduced pressure, and the resulting product purified by column chromatography on silica gel using toluene as the eluent. The benzyl protecting group was then removed by hydrogenation using a 10% Pd on carbon catalyst to give 5-hydroxy-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-
30 tetrafluoropropoxy)hexyl)-phenyl]pyrimidine.

Preparation of R-5-(5-fluoroheptyloxy)-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

To a 100 ml round bottom flask was added a 1:1 solution of dimethylformamide and acetonitrile (50 ml), potassium carbonate (0.7g, 5.1mmol), 5-hydroxy-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine (2.2g, 4.24 mmol) and R-5-fluoro-1-methylsulfonyl pentane (0.9g, 4.24mmol). The reaction mixture was heated to 80°C for 16 hours and then cooled to room temperature. The solution was diluted with toluene (100ml) and washed with water. The resulting organic phase was concentrated under reduced pressure, recrystallized from hexanes at -20°C and further purified by column chromatography eluting with 10:1 v/v hexanes/ethyl acetate to yield 2.6g of the desired product.

Example 2

Preparation of R-5-(5-fluorooctyloxy)-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

This compound was prepared essentially as described in Example 1 except methyl lithium was replaced by ethyl magnesium bromide.

Example 3

Preparation of R-5-(6-fluoroheptyloxy)-2-[4-(6-(3-nonafluorobutoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

This compound was prepared essentially as described in Example 1 except methyl lithium was replaced by 4-benzyloxybutyl-1-magnesium bromide, the intermediate epoxide was reduced with lithium triethyl borohydride and 3-perfluorobutoxy-2,2,3,3-tetrafluoropropanol was used in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol.

Example 4

Preparation of 3-Fluoro-1-methylsulfonyl hexane

This intermediate was prepared essentially as described in Example 1 using ethyl magnesium bromide and R-4-benzyloxy-1,2-epoxybutane (prepared essentially as described in J.A. Frick, Synthesis, vol. 7, pp. 621-3 (1992))

Preparation of R-5-(3-fluorohexyloxy)-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine

5 A 50 ml round bottom flask was charged with 5-hydroxy-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine (2.0g, 2.52mmol prepared as described in U.S. Patent 5,474,705), potassium carbonate (0.42g, 3.0 mmol), dimethyl formamide (7.8 ml) and 3-fluoro-1-methylsulfonyl hexane (0.55g, 2.77 mmol). The reaction mixture was heated to 100°C for 1 hour, then cooled to room temperature. The solution was diluted with 20 ml of toluene and then washed with water. The organic phase was isolated and concentrated under reduced pressure. The crude product was further purified by recrystallization from heptane, followed by and chromatography, eluting with 10:1 v/v hexane/ethyl acetate to give 1.13g of the desired product.

15

Example 5

Preparation of (R) 5-(4-fluorodecyloxy)-2-[4-(6-(2-pentafluoroethoxy-2,2-difluoroethoxy)hexyl)-phenyl]pyrimidine

The title compound was prepared essentially as described in Example 1. R-4-fluoro-1-methanesulfonyldecane was prepared by copper catalyzed addition of vinyl 20 magnesium bromide to epichlorohydrin to yield 5-chloro-4-hydroxypent-1-ene which was treated with base to give the corresponding epoxide. The epoxide is then ring opened by the addition of propyl magnesium bromide to yield 4-hydroxy-dec-1-ene. This hydroxy compound is converted to 4-fluoro-dec-1-ene as described in Example 1, followed by 25 treatment with borane/tetrahydrofuran and hydrogen peroxide to give 4-fluoro-decanol. This alcohol was treated with methane sulfonyl chloride in the presence of triethylamine to yield the desired intermediate R-4-fluoro-1-methylsulfonyl decane.

Example 6

30 **Preparation of R-5-(6-fluorooctyloxy)-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine**

The title compound was prepared essentially as described in Example 4 using R-6-fluoro-1-methylsulfonyloxy octane (prepared from 7-benzyloxy-pent-1-ene oxide as described in Example 1) in place of R-3-fluoro-1-methylsulfonyloxyhexane.

5 **Example 7**

Preparation of S-5-(6-fluoro-7-methoxy-heptyl)-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine

10 The title compound was prepared essentially as described in Example 1 using 5-nonafluorobutoxysulfonyloxy-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine (prepared as described in Example 1) and S-6-fluoro-7-methoxy-hept-1-ene (prepared by addition of methanol to R-1,2-epoxyhept-1-ene and subsequent fluorination as described in Example 1).

15

Example 8

Preparation of R-5-(3-fluorohexyloxy)-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

20 The title compound was prepared essentially as described in Example 1 using 3-fluoro-1-methansulfonyloxyhexane (prepared as described in Example 4) in place of 5-fluoro-1-methanesulfonyloxy-heptane.

Example 9

25 **Preparation of R-5-(3-fluorohexyloxy)-2-[4-(6-(3-nonafluorobutoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine**

30 The title compound was prepared essentially as described in Example 1 using 3-fluoro-1-methansulfonyloxyhexane (prepared as described in Example 4) in place of 5-fluoro-1-methanesulfonyloxy-heptane and 3-nonafluorobutoxy-2,2,3,3-tetrafluoropropanol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol (prepared as described in Example 1).

Example 10

Preparation of R-5-(3-fluorohexyloxy)-2-[4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-3-fluorophenyl]pyrimidine

5 The title compound was prepared essentially as described in Example 4 with the use of 5-benzyloxy-2-[3-fluoro-4-hydroxyphenyl]pyrimidine (prepared from 3-fluoro-4-hydroxybenzamidinium hydrochloride and $(\text{CH}_3)_2\text{NCH}=(\text{OCH}_2\text{C}_6\text{H}_5)\text{CHO}$).

Example 11

10 **Preparation of R-5-(3-fluorooctyloxy)-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine**

 The title compound was prepared essentially as described in Example 1 using 3-fluoro-1-methanesulfonyloctane (prepared as described in Example 4 using butylmagnesium bromide) in place of 5-fluoro-1-methanesulfonyl-heptane.

15

Example 12

Preparation of R-5-(3-fluorohexyloxy)-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

 The title compound was prepared essentially as described in Example 1 using 3-fluoro-1-methanesulfonyloctane (prepared as described in Example 4 using butylmagnesium bromide) in place of 5-fluoro-1-methanesulfonyl-heptane and 3-nonafluorobutoxy-2,2,3,3-tetrafluoropropanol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol (prepared as described in Example 1).

20

Examples 13

Preparation of R-5-(5-fluorohexyloxy)-2-[4-(6-(2-(2-(2-pentafluoroethoxy)-2,2,3,3-tetrafluoroethoxy)-2,2-difluoroethoxy)hexyl)-phenyl]pyrimidine

 The title compound was prepared essentially as described in Example 1 using 5-fluoro-1-methanesulfonylhexane in place of 5-fluoro-1-methanesulfonyl-heptane and 2-(2-(2-pentafluoroethoxy)-2,2,3,3-tetrafluoroethoxy)-2,2-difluoroethan-1-ol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol.

30

Example 14

Preparation of R-5-(5-fluorohexyloxy)-2-[4-(6-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

5 The title compound was prepared essentially as described in Example 1 using 5-fluoro-1-methanesulfonyhexane in place of 5-fluoro-1-methanesulfony-heptane

Example 15

10 **Preparation of R-5-(5-fluorohexyloxy)-2-[4-(6-(3-nonafluorobutoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine**

 The title compound was prepared essentially as described in Example 3 using 5-fluoro-1-methanesulfonyhexane in place of 6-fluoro-1-methanesulfony-heptane

Example 16

15 **Preparation of R-5-(5-fluoroheptyloxy)-2-[4-(6-(2-(2-(2-pentafluoroethoxy)-2,2,3,3-tetrafluoroethoxy)-2,2-difluoroethoxy)hexyl)-phenyl]pyrimidine**

 The title compound was prepared essentially as described in Example 1 using 2-(2-(2-pentafluoroethoxy)-2,2,3,3-tetrafluoroethoxy)-2,2-difluoroethan-1-ol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol.

20

Example 17

Preparation of R-5-(5-fluoroheptyloxy)-2-[4-(6-(3-nonafluorobutoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

25 The title compound was prepared essentially as described in Example 1 using 3-perfluorobutoxy-2,2,3,3-tetrafluoropropanol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol.

Example 18

30 **Preparation of R-5-(5-fluorooctyloxy)-2-[4-(6-(2-(2-(2-pentafluoroethoxy)-2,2,3,3-tetrafluoroethoxy)-2,2-difluoroethoxy)hexyl)-phenyl]pyrimidine**

The title compound was prepared essentially as described in Example 2 using 2-(2-(2-pentafluoroethoxy)-2,2,3,3-tetrafluoroethoxy)-2,2-difluoroethan-1-ol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol.

5 Example 19

Preparation of R-5-(5-fluoroheptyloxy)-2-[4-(6-(3-nonafluorobutoxy-2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

The title compound was prepared essentially as described in Example 2 using 3-perfluorobutoxy-2,2,3,3-tetrafluoropropanol in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropanol.

Example 20

Preparation of S-5-(6-fluorooctyl)-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine

The title compound was prepared essentially as described in Example 7 using R-6-fluorooct-1-ene in place of S-6-fluoro-7-methoxy-hept-1-ene.

Example 21

Preparation of S-5-(6-fluorooctyl)-2-[4-(2-(2-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2-tetrafluoroethoxy)2,2-difluoroethoxy)-phenyl]pyrimidine

The title compound was prepared essentially as described in Example 7 using R-6-fluorooct-1-ene in place of S-6-fluoro-7-methoxy-hept-1-ene and 5-nonafluorobutoxysulfonyl-2-[4-(2-(2-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2-tetrafluoroethoxy)2,2-difluoroethoxy)-phenyl]pyrimidine in place of 5-nonafluorobutoxysulfonyl-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-phenyl]pyrimidine.

Example 22

Preparation of S-5-(6-fluorooctyl)-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-3-fluorophenyl]pyrimidine

The title compound was prepared essentially as described in Example 20 using 5-benzyloxy-2-[3-fluoro-4-hydroxyphenyl]pyrimidine (prepared from 3-fluoro-4-hydroxybenzamidinium hydrochloride and $(\text{CH}_3)_2\text{NCH}=(\text{OCH}_2\text{C}_6\text{H}_5)\text{CHO}$) in place of 5-benzyloxy-2-[4-hydroxyphenyl]pyrimidine.

5

Example 23

Preparation of R-5-(6-fluorooctyloxy)-2-[4-(3-(2-(2-(2-trifluoromethoxy)-1,1,2,2-tetrafluoroethoxy)-1,1,2,2-tetrafluoroethoxy) 2,2-difluoroethoxy)propyl)-phenyl]pyrimidine

10

The title compound was prepared essentially as described in Example 1 with the use of R-6-fluoromethylsulfonyl octane in place of 5-Fluoro-1-methylsulfonyl heptane and 2-(2-(2-trifluoromethoxy)-1,1,2,2-tetrafluoroethoxy)-1,1,2,2-tetrafluoroethoxy 2,2-difluoroethoxy)prop-3-ene in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene

15

Example 24

Preparation of R-5-(6-fluorooctyloxy)-2-[4-(6-(3-pentafluoroethoxy)- 2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

20

The title compound was prepared essentially as described in Example 1 with the use of R-6-fluoromethylsulfonyl octane in place of 5-Fluoro-1-methylsulfonyl heptane.

Example 25

Preparation of R-5-(6-fluorooctyloxy)-2-[4-(6-(3-perfluorobutoxy)- 2,2,3,3-tetrafluoropropoxy)hexyl)-phenyl]pyrimidine

25

The title compound was prepared essentially as described in Example 1 with the use of R-6-fluoromethylsulfonyl octane in place of 5-Fluoro-1-methylsulfonyl heptane 3-perfluorobutoxy- 2,2,3,3-tetrafluoropropoxyhex-6-ene in place of 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene.

30

Example 26

Preparation of R-5-(6-fluorooctyloxy)-2-[4-(4-(4-(1,1,2,2,3,3,4,4,4-nonafluorobutoxy)-1,1,2,2,3,3,4,4-octafluorobutoxy)2,2,3,3,4,4-hexafluorobutoxy)-3-fluorophenyl]pyrimidine

The title compound was prepared essentially as described in Example 6 using 5-benzyloxy-2-[3-fluoro-4-hydroxyphenyl]pyrimidine (prepared from 3-fluoro-4-hydroxybenzamide hydrobromide and $(\text{CH}_3)_2\text{NCH}=\text{C}(\text{OC}_8\text{H}_{17})\text{CHO}$).

Preparation of Comparative Examples C-1 through C-6

Comparative Example 1

The compound was prepared essentially as described in Example 1 by combining 5-hydroxy-2-[4-(6-(2-(2-trifluoromethoxy)-tetrafluoroethoxy)-2,2,-difluoroethoxy)hexyl)-phenyl]pyrimidine (prepared essentially as described in Example 1 using 1-(2-(2-trifluoromethoxy)-tetrafluoroethoxy)-2,2,-difluoroethoxyhex-6-ene in place of 1-(3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene) and R-2-fluoro-1-methylsulfonyl octane (prepared from R-2-fluorooctan-1-ol and methylsulfonyl chloride) in place of 5-fluoro-1-methylsulfonyl heptane.

Comparative Examples 2 and 3

These compounds were prepared essentially as described in Comparative Example 4 and Example 5, respectively, of U.S. Patent No. 5,702,637.

Comparative Example 4

(R) 5-Octyloxy-2-[4(7-(3-nonafluorobutoxy)-2,2,3,3-tetrafluoropropoxy)-6-fluoroheptyl]phenyl]pyrimidine

The compound was prepared essentially as described in Example 1 with the substitution of R-(3-nonafluorobutoxy)-2,2,3,3-tetrafluoropropoxy 6-fluorohept-1-ene for 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene and substitution of 5-octyloxy-2-[4-hydroxyphenyl]pyrimidine] for 5-benzyloxy-2-[4-hydroxyphenyl]pyrimidine. R-(3-nonafluorobutoxy)-2,2,3,3-tetrafluoropropoxy 6-fluorohept-1-ene can be prepared by

reaction of 1-butenyl-4-magnesium bromide and epichlorohydrin followed by addition of 3-nonafluorobutoxy-2,2,3,3-tetrafluoropropanol and subsequent fluorination as described in Example 1

5 Comparative Example 5

(R) 5-Hexyloxy-2-[4(7-(3-nonafluorobutoxy)-2,2,3,3-tetrafluoropropoxy)-7-fluorooctyl)phenyl]pyrimidine

The compound was be prepared essentially as described in Example 1 with the substitution of R-(3-nonafluorobutoxy)-2,2,3,3-tetrafluoropropoxy -7-fluorooct-1-ene for
 10 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene and substitution of 5-hexyloxy-2-[4-hydroxyphenyl]pyrimidine] for 5-benzyloxy-2-[4-hydroxyphenyl]pyrimidine. R-(3-nonafluorobutoxy)-2,2,3,3-tetrafluoropropoxy 7-fluorooct-1-ene can be prepared by reaction of 1-pentenyl-5-magnesium bromide and epichlorohydrin followed by addition of
 15 in Example 1.

Comparative Example 6

(R) 5-Heptyloxy-2-[4(7-(3-nonafluorobutoxy)-2,2-difluoroethoxy)-7-fluorooctyl)phenyl]pyrimidine

The compound was prepared essentially as described in example 1 with the substitution of R-(3-nonafluorobutoxy)-2,2-difluoroethoxy -7-fluorooct-1-ene for 3-pentafluoroethoxy-2,2,3,3-tetrafluoropropoxy)hex-6-ene and substitution of 5-heptyloxy-2-[4-hydroxyphenyl]pyrimidine] for 5-benzyloxy-2-[4-hydroxyphenyl]pyrimidine. R-(3-nonafluorobutoxy)-2,2-difluoroethoxy 7-fluorooct-1-ene can be prepared by reaction of 1-
 25 pentenyl-5-magnesium bromide and epichlorohydrin followed by addition of 3-nonafluorobutoxy-2,2-difluoroethanol and subsequent fluorination as described in Example 1.

Table 1

Mesomorphic Phase Transitions (°C)

Exempl e	Structure	I - Sa	to Sc	to Sm	to K	to Sc
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Table 1
Mesomorphic Phase Transitions (°C)

Exempl e	Structure	I - Sa	to Sc	to Sm	to K	to Sc
1		96	57	9	-10	18
2		98	73	1	-9	9
3		105	79	-11		-8
4		141	89	43		49
5		81	72	17		32
6		130	97	53		63
7		111	72	10		31
8		91.5		-31		-10
9		100	50	-25		-14
10		123	78	36.2		43.3
11		95	64	-3		10
12		102	78	0		10
13		103	49	-7	-22	5
14		96	22	-7		5
15		103	57	-4		7
16		102	69	6	-15	15

Table 1

Mesomorphic Phase Transitions (°C)

Exempl e	Structure	I - Sa	to Sc	to Sm	to K	to Sc
17		103	73	10		18
18		104	82	1.3		9
19		104	84	0		5
20		114	57	39	-37	48
21		84	46	27		44
22		93	52	24		36
23		118	83	-11		-1
24		97	79	-8		1
25		105	88	-7		2
26		107	88	56		66
C-1		I to Sc	68.7	44.2	11.9	52.8
C-2		99	71	60		73
C-3		91	64	-7		43
C-4		105	63	-15		22
C-5		97	68	-3		14
C-6		91	67	1		13

The data of Table 1 shows that the compounds of the invention have broader mesomorphic temperature ranges for device use from -10° to 80°C and are more stable in these phases at lower temperatures than the compounds of Comparative Examples 1 and 2. In comparison with the compounds of Comparative Examples 3-6, the compounds of the invention have similar mesomorphic temperature ranges for device use from -10° to 80°C .

Examples 27 to 32 and Comparative Examples C-7 to C-11

A series of devices, each containing a chiral compound of this invention or a comparative compound, were prepared essentially as described in U.S. Patent No. 5,377,033 (Radcliffe). The ITO-constituted electrodes of each device were connected to a function generator with variable output voltage. The device was first heated to the isotropic temperatures and then cooled to the reduced temperature ($T-T_{A-C}$, the temperature below the SmA to SmC transition temperature) noted in Table 2 (below). The polarization (nC/cm^2) and the τ_{electric} were determined as described below:

The polarization of the devices were determined essentially as described by Miyasato et al. in Jap. J. Appl. Phys. 22, 661 (1983). The electronic response time, τ_{electric} , was derived from the displacement current of the ferroelectric liquid crystal device under an applied square voltage pulse. The current was viewed on a 100 megahertz bandwidth oscilloscope. The usual decaying exponential, associated with a dielectric filled capacitor, was followed by the spontaneous polarization (P_s) switching pulse. The time from the rising edge of the voltage pulse to the peak of the P_s pulse was taken to be τ_{electric} . In these Examples subphases of the chiral tilted smectic mesophases, such as antiferroelectric or ferrielectric phases, were not specified.

Table 2				
Example	Compound from:	$T-T_{A-C}$ ($^{\circ}\text{C}$)	P_s (nC/cm^2)	TAU (μsec)
27a	Ex 1	-7	13.4	11.1

Table 2				
Example	Compound from:	T-T _{A-C} (°C)	Ps (nC/cm ²)	TAU (μsec)
27b	"	-37	29.9	25.7
28a	Ex 2	-22	13.4	15
29a	Ex 3	-28	26.8	11.7
29a	"	-58	45.8	25.3
30a	Ex 4	-10	24.4	8.8
30a	"	-39	48	17.5
31a	Ex 5	-11	-14.2	11.1
31b	"	-36	18	28.7
32a	Ex 8	-10	*	18.1
32b	"	-20	*	25.7
C-7a	C-1	-10	89	7.5
C-7b	"	-20	113	8.9
C-8a	C-2	-4	53	10.4
C-9a	C-3	-14	103	5.0
C-9b	"	-34	139	8.8
C-10a	C-4	-20	14	14
C-10b	"	-40	18	27
C-11a	C-5	-20	26	8
C-11b	"	-40	38	13

*Unable to measure under standard conditions

The results in Table 2 show that the compounds of Comparative Examples C-7, C-8 and C-9 had large polarization values (typically greater than 100 nC/cm² below T_{A-C} - 20°) in comparison to the compounds of the invention (typically less than 50 nC/cm² below T_{A-C} - 20°). Low polarization values are important for the practical operation of a liquid crystal device. Polarization reversal fields, which cause switching or partial switching back to a material's original director alignment, are larger for higher polarization liquid crystal compounds or mixtures. This polarization reversal results in loss of the bistability that is crucial to the passive-matrix driving of ferroelectric liquid crystal devices. Another potential disadvantage of using high polarization mixtures is the partial switching of director alignment in response to non-switching (secondary) signals in a driving waveform. This continued response or fluctuation of the director causes a large decrease in the contrast ratio of a ferroelectric liquid crystal device. This indicates that these comparative materials should be diluted with achiral materials in order to reduce the polarization of the mixtures

The compounds of the invention show similar polarization values and response times in comparison with the compounds of Comparative Examples C-10 and C-11. However, the compounds of the invention allow much easier synthetic variation of the terminal portion comprising the extended group, than the compounds of the Comparative Examples (where the compounds have a chiral, extended fluorochemical terminal). Structural changes in this portion of the molecule have large effects on mesomorphic transition temperatures, tilt angles, and other properties. The benefits and effects of changing this portion of the molecule have been described Applicant's U.S.S.N. 08/827,753 and U.S.S.N. 08/827,287.

Further, as can be seen in Table 2, the compounds of this invention had a high temperature dependence of the polarization P_S. An increase in the temperature dependence of the polarization results in a decrease in the temperature dependence of τ_{electric} .

Surprisingly, Example 31a,b of Table 2 exhibited a polarization behavior (nC/cm²) that inverted with a decrease in temperature (°C) from the Smectic A mesophase to the Smectic C mesophase. Figure 1 (below) shows the change in polarization for Examples 27a,b and 31a,b of the invention and demonstrates a change from a negative to a positive

polarization value for Example 31 of the current invention. This enables the formulation of mixtures with temperature independent switching as shown in Example 36(below).

Examples 33 to 35 and Comparative Examples C-12 to C-13

5 When the τ_{electric} (switching time) and temperature data in Table 2 is plotted as the $\ln(\tau_{\text{electric}})$ versus temperature, relatively small changes in response times are observed over the temperature ranges, demonstrating that the liquid crystal compounds of the invention have the desirable property of reduced temperature dependent switching; this data is tabulated below in Table 4, as the slope (change in $\ln(\tau_{\text{electric}})$)/change in
10 temperature) along with the average polarization and average temperature of the range. For comparative purposes, examples from U. S. Patent Nos. 5,262,388 and 5,702,637 are included in Table 4. In Table 4 Tc-10 to Tc-40 refers to the temperature range from -10°C to -40°C below the Smectic A to Smectic C transition temperature. Comparative Example
15 13 is a mixture disclosed in Example 36 of U. S. Pat. 5,702,637 (Johnson et al.). Shown below in Tables 3 and 4 are the compounds used and the weight percents used in preparing the mixture of this Comparative Example.

Table 3

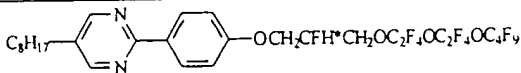
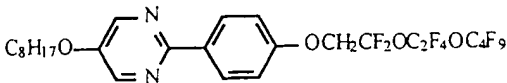
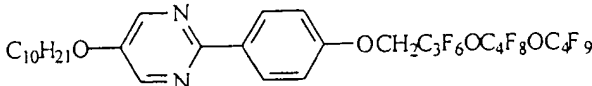
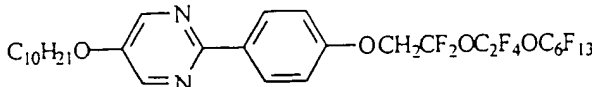
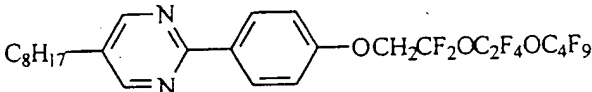
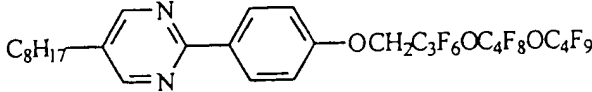
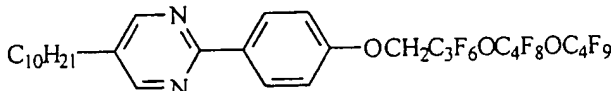
Example	Compound	Reference	Weight %
C-13		Ex 36 US 5702637	9.7
			11.5
			11.6
			12.6
			6.6
			13.4
			34.7

Table 4
Temperature dependence of switching
over temperature range of T_c-10 to T_c-40

Example	Compound from:	Slope	average P _s (nC/cm ²)	average Temp (°C)
33	Ex 1	-.028	22	35
34	Ex 2	-.017	17	44
35	Ex 3	-.018	24	56
C-12	C-3	-.029	127	35
C-13	Table 3	-.038	11	45

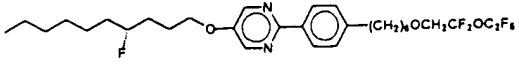
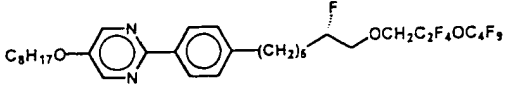
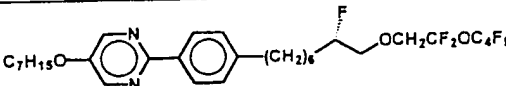
The data in Table 4 shows that similar, and in some cases, less temperature dependant-switching behavior was obtained with compounds of the invention relative to the compounds and composition of Comparative Examples C-12 and C-13. The compounds of Comparative Example C-13 illustrates the effect on the temperature dependence of switching by diluting a high polarization material with achiral compounds. In this comparison, Comparative Example C-12 provided good temperature dependence of switching when used at 100 wt.%, but it had a polarization that was too high for some device applications. Upon diluting this compound with achiral materials, the polarization was lowered to the range of compounds of the invention, however, the temperature dependency of switching increased dramatically.

Example 36

In the following Example, a device was prepared as described in Example 27 and filled with the mixture shown in Table 5. The temperature dependence of Tau (τ_{electric} switching time in μsec) was evaluated from T_{A-C} -10°C to T_{A-C} -40°C (the temperature range below the Smectic A to Smectic C transition temperature) by analysis of the change in $(\ln \tau) / \Delta T$ (in degrees Centigrade below the

Smectic A to Smectic C transition). Shown below in Table 5 are the compounds and the weight percents used to prepare the mixture of this Example along with the slope values. Also in Figure 2 (below) the polarization vs. temperature of the mixture is plotted for the mixture and its constituent compounds. In Table 2 polarization values are in nC/cm^2 and temperature is in $^{\circ}\text{C}$.

Table 5

Example	Compound	Reference	Weight %	Tau slope
36		Ex 5	50	-0.007
		C-4	25	
		C-6	25	

As can be seen in Table 5 and Figure 2, the mixing of a material which has a negative to positive polarization inversion effect (on cooling) with materials which have positive polarization led to a mixture with a decreased temperature dependence of switching (Figure 3). Conversely the mixing of a material which has a positive to negative polarization inversion effect (on cooling) with materials which have negative polarization can also lead to a mixture with a decreased temperature dependence of switching. In these cases the resultant mixture has an increased temperature dependence of polarization, which leads to a decreased temperature dependence of the switching properties.

When the tau electric (switching time) and temperature data in Table 2 is plotted as the teletric versus temperature as shown in Figure 3, relatively small changes in response times are observed over the temperature ranges, demonstrating that the liquid crystal materials of the this invention have the desirable property of reduced temperature dependent switching; this data is tabulated above in Table 4, as the slope (change in $\ln(\text{tau electric}) / (\text{change in temperature})$) along with the average polarization and average temperature of the range. For comparative purposes examples from U. S. Patents

5,262,388 and 5,702,637 are included in Table 3. Comparative Example 13 is a mixture disclosed in Example 36 of U. S. Pat. 5,702,637 (Johnson et al.).

Example 37

5 A cell of the type described in U.S. 5,377,033 (Radcliffe) was filled with the compound of Example 6 and placed in a hot stage with openings to allow transmitted light. The hot stage and the cell were placed between the crossed polarizers of a transmitting, polarizing microscope. The microscope was equipped with a HAMAMATSU Model HC
10 124-01 photomultiplier tube (PMT) and amplifier to detect the transmitted light levels. The PMT amplifier output was connected to a TEKTRONIX Model TDS 420 oscilloscope. A triangle waveform test signal was generated by a WAVETEK Model 395 arbitrary function generator. The signal from the generator was amplified through a Krone-Hite Model 7602 wideband amplifier. The oscilloscope was set to plot the
15 transmission signal versus the triangle waveform test signal. The cell was heated to about 10 degrees above the isotropic transition of the liquid crystal composition and was cooled back down at 0.5 degrees per minute to the smectic A phase, without any signal being applied to the cell.

The microscope stage was rotated to align a zero field optical axis (smectic layer
20 normal) of the liquid crystal composition with one of the crossed polarizers. The cell was cooled to a few degrees above a tilted smectic phase of the liquid crystal composition. To enable the tilted phase to be detected, the triangle signal was applied to the cell. A phase change (at T_{A-C}) from the smectic A mesophase to the tilted smectic mesophase was detected by a change in transmission signal to a non-linear response. In a smectic A
25 mesophase, any transmission signal from the PMT is due to an electroclinic effect which gives a domainless, linear transmission response. In a tilted smectic or induced tilted smectic mesophase, the transmission response becomes non-linear. Transmission versus voltage curves were generated at selected frequencies and selected temperatures, (80°C and 90°C, indicated in Figures 4 and 5 below, respectively).

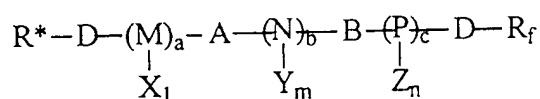
As can be seen in Figures 4 and 5, the compound of this Example exhibited tristable switching, a distinct threshold, a double hysteresis allowing for a memory effect in either of the driven states, and was easily switched to produce a device having few defects and allowing for the recovery of alignment. The micro-second switching speed
5 observed for this compound is considerably faster than the millisecond switching speed of currently used nematic liquid crystal compounds.

Various modifications and alterations of this invention will become apparent to those skilled in the art without departing from the scope and spirit of this invention.

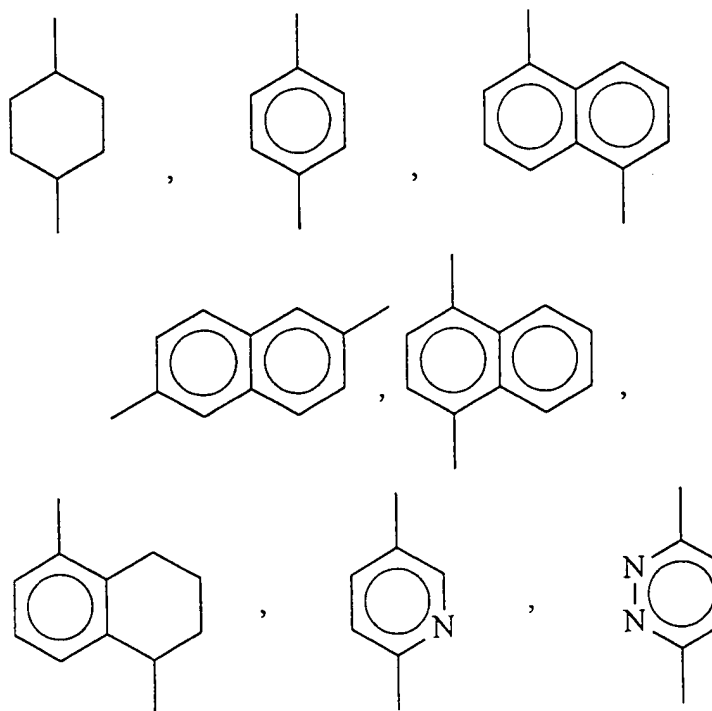
What Is Claimed Is:

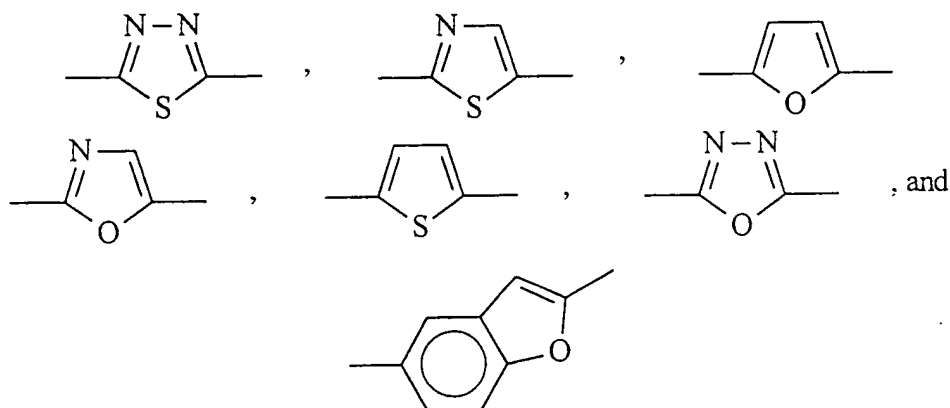
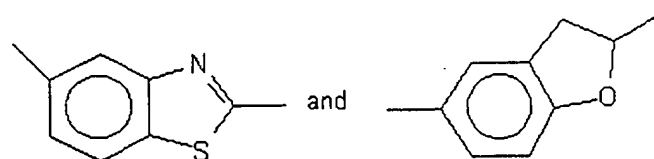
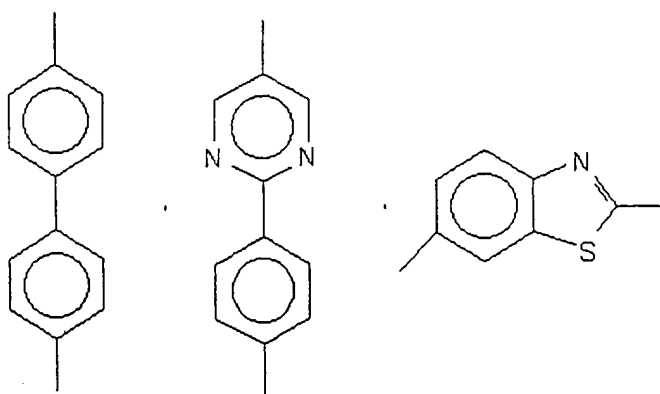
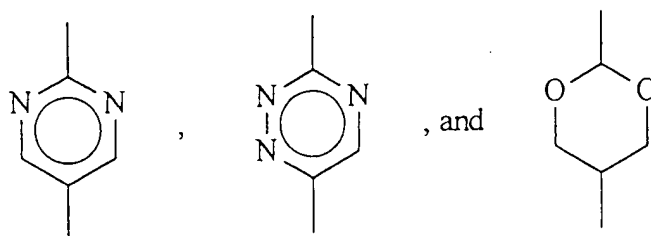
1. Fluorine containing, chiral liquid crystal compounds comprising (a) an achiral fluorochemical terminal portion which comprises a terminal fluoroalkyl, fluoroether, perfluoroalkyl, or perfluoroether group; (b) a chiral terminal portion comprising a saturated or unsaturated chiral hydrocarbon or chiral hydrocarbon ether group comprising a chiral center; and (c) a central core connecting said terminal portions; said chiral terminal portion having at least 3 in-chain atoms between said chiral center of said chiral terminal portion and said central core.

2. The compounds of Claim 1, wherein said compounds are represented by the general formula:



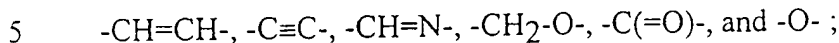
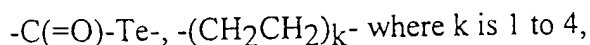
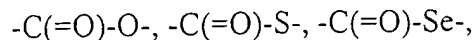
where M, N, and P are each independently selected from the group consisting of





a, b, and c are each independently zero or an integer of from 1 to 3, with the proviso that the sum of $a + b + c$ be at least 1;

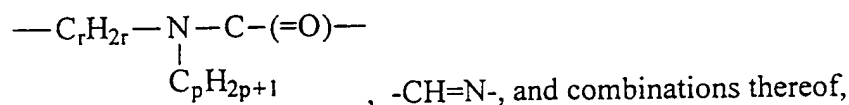
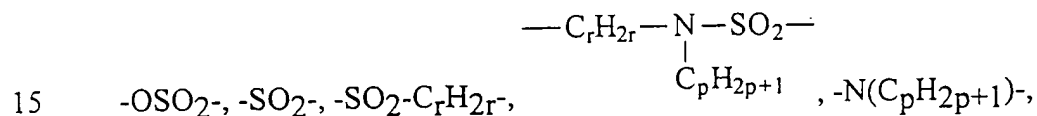
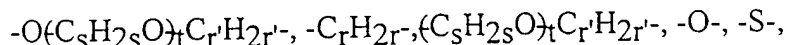
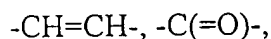
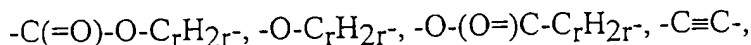
each A and B are non-directionally and independently selected from the group consisting of a covalent bond,



each X, Y, and Z are independently selected from the group consisting of -H, -Cl, -F, -Br, -I, -OH, -OCH₃, -CH₃, -CF₃, -OCF₃, -CN, and -NO₂;

each l, m, and n are independently zero or an integer of 1 to 4;

10 each D is non-directionally and independently selected from the group consisting of a covalent bond,



20 where r and r' are independently integers of 0 to about 20, s is independently an integer of 1 to about 10 for each (C₅H₂SO), t is an integer of 1 to about 6, and p is an integer of 0 to about 4;

R* is a chiral hydrocarbon or hydrocarbon ether moiety containing at least one chiral center

(asymmetric carbon atom) with the proviso that at least one chiral center of R* is spaced from the central core by at least 3 in-chain atoms;

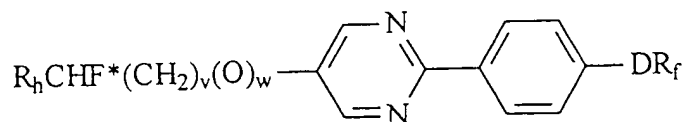
25 R_f is fluoroalkyl, perfluoroalkyl, fluoroether, or perfluoroether.

3. The compounds of Claim 2 wherein said R_f group is represented by the formula -C_qF_{2q}X', where q is an integer of 1 to about 20 and X' is hydrogen or fluorine.

4. The compounds of Claim 2 wherein R_f is represented by the formula -
 $(C_xF_{2x}O)_zC_yF_{2y+1}$, where x is independently an integer of 1 to about 12 for each
 $(C_xF_{2x}O)$, y is an integer of 1 to about 12, and z is an integer of 1 to about 10.

5

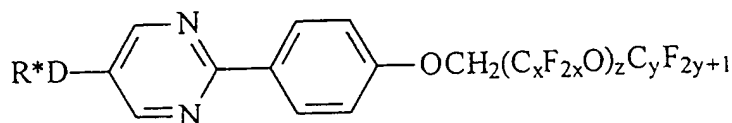
5. The compounds of Claim 2, wherein said compounds are represented by the
 formula:



- where D and R_f are as defined in Claim 3, R_h is a linear or branched alkyl group of 1 to
 10 about 10 carbon atoms, $-CHF^*$ represents a chiral center, v is an integer of from 2 to
 about 10, w is an integer of 0 or 1, with the proviso that $v+w$ is at least 3, where one or
 more non-adjacent $-CH_2-$ groups may be replaced with an oxygen atom and where one or
 more ring hydrogen atoms can be replaced by fluorine atoms.

15

6. The compounds of Claim 2, wherein said compounds are represented by the
 formula:



- where R^* and D are as defined in Claim 2, x is independently an integer of 1 to about 12
 for each $(C_xF_{2x}O)$, y is an integer of 1 to about 12, and z is an integer of 1 to about 10 and
 20 where one or more ring hydrogen atoms can be replaced with fluorine atoms.

7. The compounds of Claim 1 wherein the slope of the plot of $\Delta(\ln$
 $\tau_{\text{electric}})/\Delta T$ is greater than -0.03 .

8. A liquid crystal display device containing at least one compound of Claim

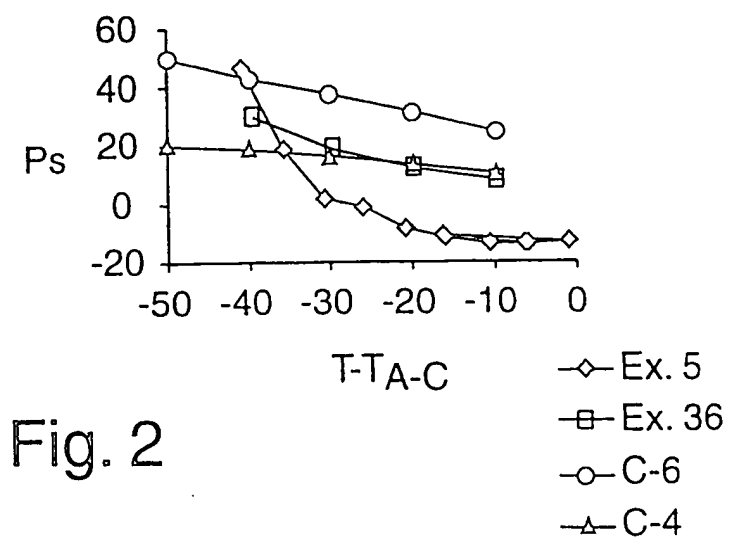
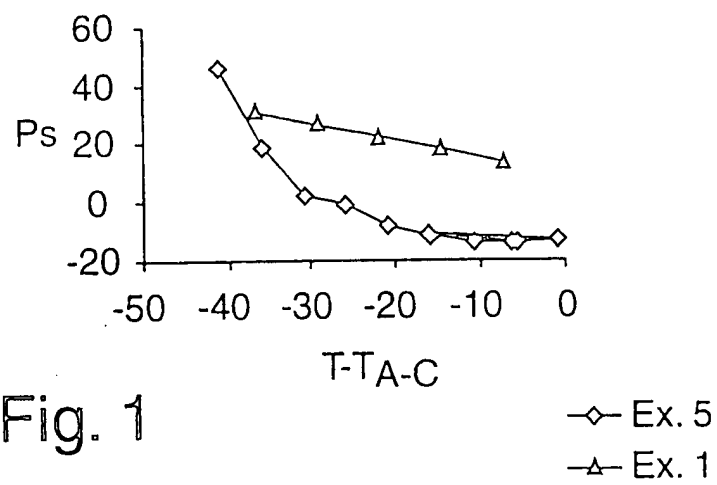
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1.

9. The device of Claim 8 wherein the polarization (P_s) value of said compound switches from positive to negative (+ to -), or negative to positive (- to +) as the temperature is reduced from the smectic A to smectic C transition temperature.

5 10. A tristable liquid crystal display device containing at least one compound of Claim 6.

1/3



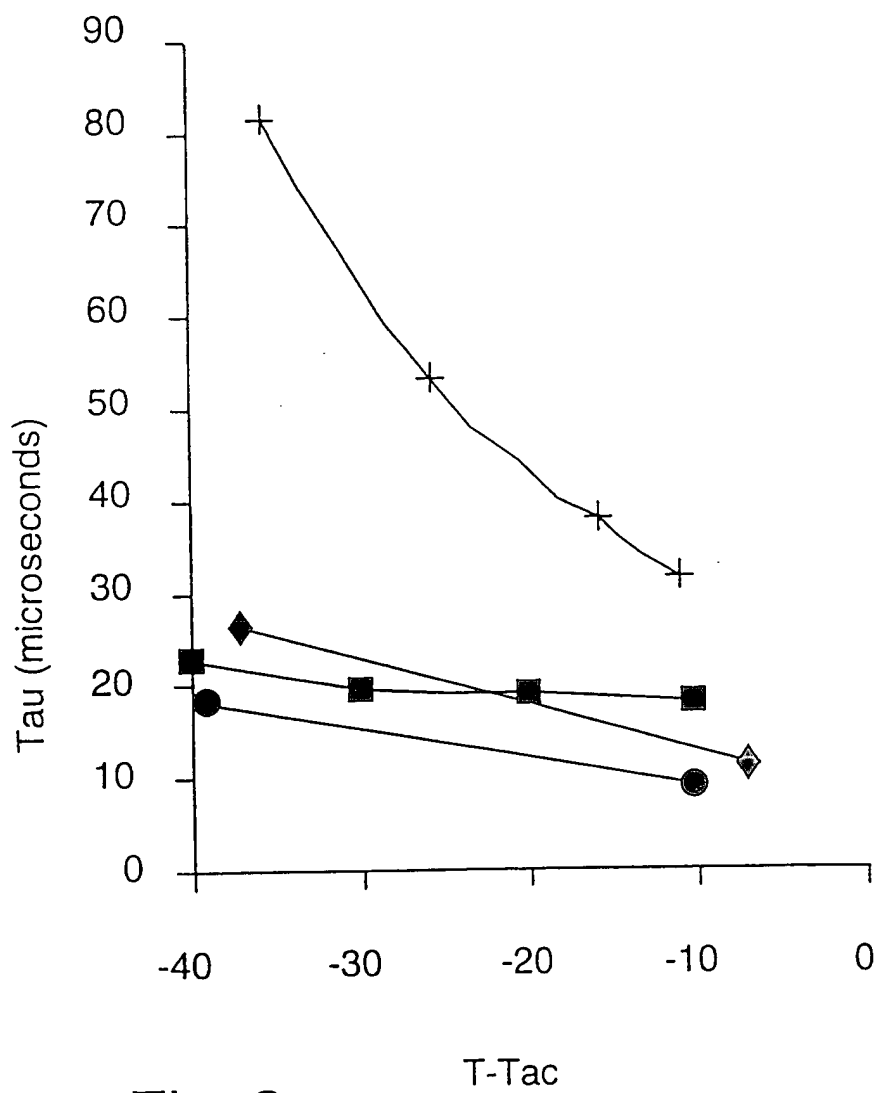
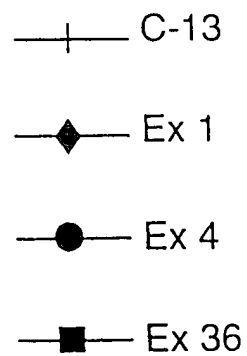
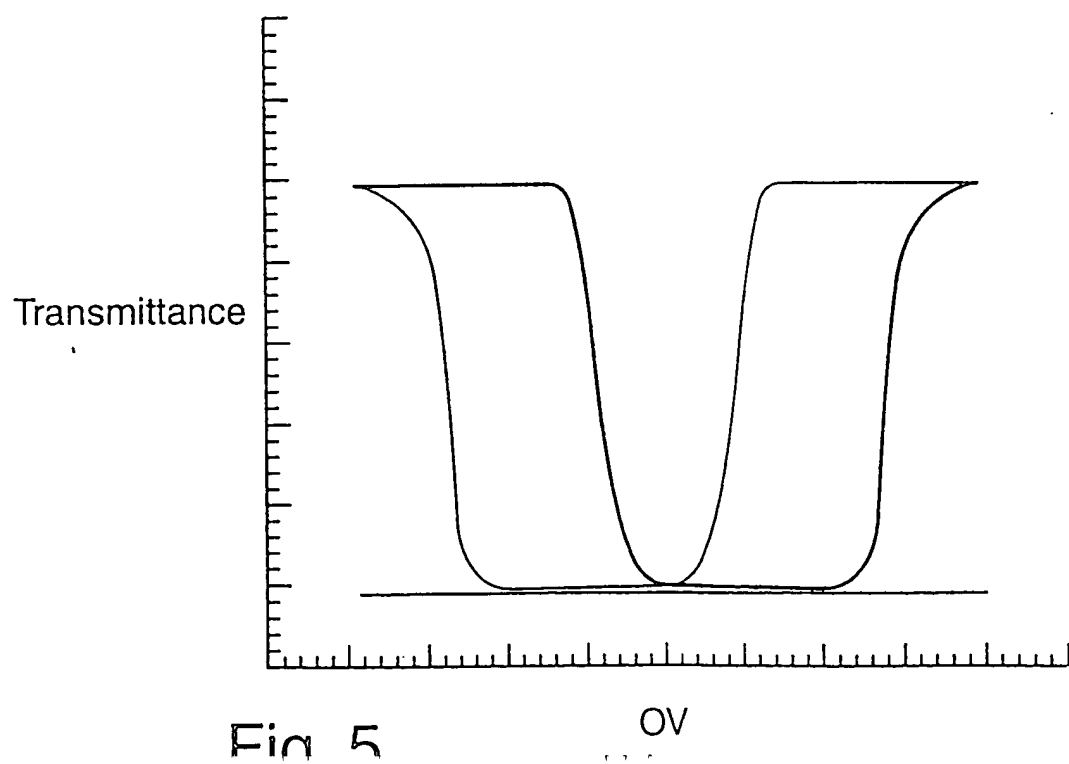
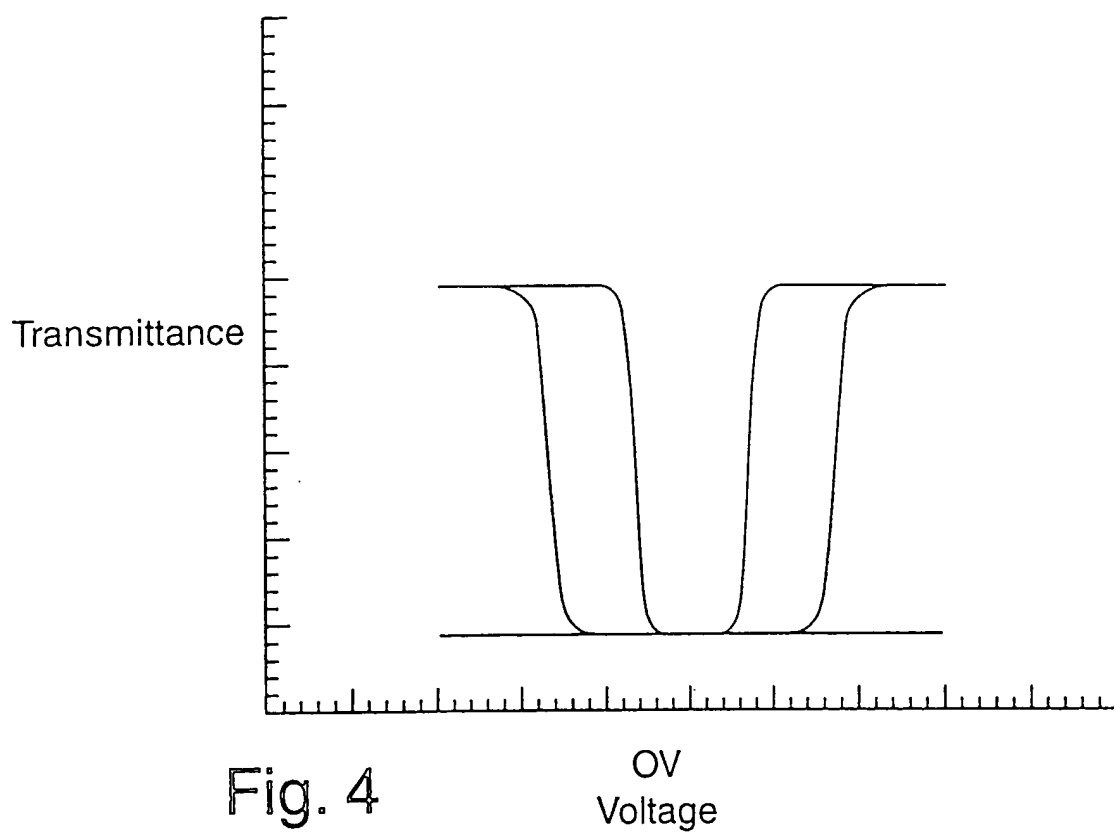


Fig. 3





INTERNATIONAL SEARCH REPORT

Internat'l Application No

PCT/US 99/05380

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C09K19/04 C09K19/34 C07D239/26 C07C43/225 C07C69/76
 C07D239/34 C09K19/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C09K C07D C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 95 17481 A (MINNESOTA MINING) 29 June 1995 see page 4, line 10 - page 5, line 41 see page 33 - page 34; table 1 see page 7, line 20 - page 10, line 29 ---	1-4, 6, 8
Y	EP 0 255 236 A (MINNESOTA MINING) 3 February 1988 see page 2, line 48 - page 3, line 18 see page 4, line 37 - page 6, line 15 ---	1-4, 6, 8
Y	WO 91 00897 A (MERCK PATENT) 24 January 1991 see page 1, line 1 - page 2, line 5 --- -/--	1-3



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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Date of the actual completion of the international search

18 June 1999

Date of mailing of the international search report

25/06/1999

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/05380

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 97 13822 A (MINNESOTA MINING) 17 April 1997 see page 4, line 22 - page 9, line 30 & US 5 658 491 A cited in the application ---	1-8
Y	US 5 702 637 A (G.C.JOHNSON) 30 December 1997 cited in the application see column 1, line 47 - column 2, line 57 see column 3, line 16 - column 7, line 6 ---	1-8
Y	US 5 417 883 A (K.A.EPSTEIN) 23 May 1995 cited in the application see column 4, line 17 - column 7, line 7 -----	1-8

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/05380

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9517481 A	29-06-1995	AU 690944 B	07-05-1998
		AU 1180595 A	10-07-1995
		CA 2177590 A	29-06-1995
		DE 69411286 D	30-07-1998
		DE 69411286 T	05-11-1998
		EP 0736078 A	09-10-1996
		ES 2117847 T	16-08-1998
		FI 962535 A	18-06-1996
		JP 9507059 T	15-07-1997
		US 5474705 A	12-12-1995
EP 255236 A	03-02-1988	DE 3789740 D	09-06-1994
		DE 3789740 T	24-11-1994
		FI 872851 A, B,	31-12-1987
		JP 2086171 C	23-08-1996
		JP 7116107 B	13-12-1995
		JP 63027451 A	05-02-1988
		JP 2705714 B	28-01-1998
		JP 7324051 A	12-12-1995
		KR 9604450 B	06-04-1996
		KR 9601889 B	06-02-1996
		US 4886619 A	12-12-1989
		US 5399701 A	21-03-1995
		US 5254747 A	19-10-1993
WO 9100897 A	24-01-1991	DE 4006743 A	24-01-1991
		DE 59009154 D	29-06-1995
		EP 0482024 A	29-04-1992
WO 9713822 A	17-04-1997	US 5658491 A	19-08-1997
		EP 0868501 A	07-10-1998
US 5702637 A	30-12-1997	CA 2217608 A	24-10-1996
		EP 0821719 A	04-02-1998
		WO 9633251 A	24-10-1996
US 5417883 A	23-05-1995	CA 2185797 A	19-10-1995
		EP 0755425 A	29-01-1997
		JP 10501562 T	10-02-1998
		WO 9527765 A	19-10-1995